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510(k) Premarket Notification Database

Device Classification Name [Computer, Diagnostic, Programmable](#)

510(K) Number K992703

Device Name CARDIOTRON EKG MULTI-PHASE INFORMATION ANALYSIS S PREMIER HEART, LLC.

Applicant 601 13th Street, N.W. Suite 901 South Washington, DC 20005

Contact William D Hare

Regulation Number [870.1425](#)

Classification Product Code [DQK](#)

Date Received 08/12/1999

Decision Date 03/21/2000

Decision Substantially Equivalent (SE)

Classification Advisory Committee Cardiovascular

Review Advisory Committee Anesthesiology

Statement/Summary/Purged Status Summary/Purged 510(K)

Summary [Summary](#)

Type Traditional

Reviewed By Third Party No

Expedited Review No

Database Updated 09/08/2008